PTO/SB08a (05-07)
Approved for use through 09/30/2007 OMB 0851-0331
U.S. Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE Under the Paperwork Reduction Act of 1995, no persons are required to resp and to a collection of information unless it contains a valid OMB control number.

	Application Number		10655889		
	Filing Date		2003-09-04		
NFORMATION DISCLOSURE	First Named Inventor	Richa	ard Schmidt		
STATEMENT BY APPLICANT Not for submission under 37 CFR 1.99)	Art Unit		1657		
not for Gabiniosion and Gr Gr R 1.50)	Examiner Name Afren		remova, Vera		
	Attorney Docket Numb	er	5662-1-PUS-1-1		

					U.S.I	PATENTS			Remove		
Examiner Cite Initial* Code1 Patent Number Kind Code1 Issue Date)ate	Name of Patentee or Applicant Relev			es,Columns,Lines where want Passages or Relevant res Appear				
	1										
If you wisl	h to a	dd additional U.S. Pate	nt citatio	n inform	ation pl	ease click the	Add button.	-	Add		_
			U.S.P	ATENT	APPLIC	CATION PUBL	LICATIONS		Remove		_
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	Name of Patentee or Applicant Re		Releva	Columns,Lir int Passages s Appear		
	1										
If you wisl	h to a	dd additional U.S. Publi	shed Ap	plication	citation	n information p	lease click the Ad	d button			
				FOREIG	SN PAT	ENT DOCUM	ENTS		Remove		
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	or ,	Pages,Colun where Releva Passages or Figures Appe	ant Relevant	Τē
	1										С
If you wis	h to a	l dd additional Foreign P	atent Do	cument	citation	information pl	I lease click the Add	button	Add		_
			NON	I-PATE	NT LITE	RATURE DO	CUMENTS		Remove		_
Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, pages(s), volume-issue number(s), publisher, city and or country where published.							Τs				

	1	Medical Pharmacology at a Glance, Blackwell Scientific Publications, 1987- pp. 18-19.	
	2	1996 MIMS Annual, Australian Edition. The entry for Boltulinum Toxin is located at 5-372.	
If you wis	h to a	dd additional non-patent literature document citation information please click the Add button Add	 1

EXAMINER SIGNATURE

Date Considered

Date Considered

"EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 809. Draw line through a

citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

1 See Kind Codes of USPTO Patient Documents at <u>www.USPTO.GDV</u> or MPEP 901.04. ² Entire office that issued the document, by the two-letter code (WIPD Standard STs.). ² For Japanese peter documents, we acticate on the year of the region of the Engineer must precise the sent another of the patient document. ² Wind of documents by the paperpartate yearhood as included on the document under WIPD Standard STs.18 pound. ² Applicate to be pade a chick many network.

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number			10655889				
	Filing Date		2003-09-04				
First Named Inventor Richa			rd Schmidt				
	Art Unit		1657				
			ova, Vera				
			5662-1-PUS-1-1				

CERTIFICATION STATEMENT

Please see	37	CFR :	1 97	and	1 98 1	n make	the	appropriate	selection(s):	

That each item of information contained in the information disclosure statement was first cited in any communication
from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the
information disclosure statement. See 37 CFR 1.97(e)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office is a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1/56(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1/97(c).

See attached certification statement.	

ľ	П	Fee set forth in 37	CFR 1.17 (p)	has been submitted he	erewith.

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Daria G. Yoerg/	Date (YYYY-MM-DD)	2007-08-23
Name/Print	Darla G. Yoerg	Registration Number	48053

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life railed by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 12.0 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case: Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. operatment of Commence, P.O. 8bx 1449, Alexandriv, V.S. 2311-1450, D.O. NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. 8bx 1459, Alexandriva, V.S. 2311-1450.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the stacked form related to a petient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is civulating; and (3) the principal purpuse for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or cosmisting your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or cosmisting your submission related to a patient agricultant or patient. If you do not furnish the requested process and the process of the process and the process of the pro

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these records.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record perfains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
 application pursuant to 35 U.S.C. 12(2) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
 disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
 which became abandoned or in which the proceedings were terminated and which application is referenced by either a
 published application, an application open to public inspections or as issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.